

19. The method of claim 4, wherein each dose contains about 50 mg to about 65 mg of the neridronic acid.

20. The method of claim 19, wherein the neridronic acid is administered about every three days.

21. The method of claim 5, wherein each dose contains about 100 mg of the neridronic acid.

22. The method of claim 3, wherein each dose contains an equivalent of about 50 mg to about 60 mg of the neridronic acid in an acid form.

23. The method of claim 1, wherein the neridronic acid is administered about once daily to about once weekly.

24. The method of claim 1, wherein the neridronic acid is administered in a single or in divided doses.

25. The method of claim 17, wherein about four doses of about 100 mg of the neridronic acid are administered.

26. The method of claim 1, wherein the neridronic acid is administered weekly for about four to about six weeks.

27. The method of claim 1, wherein the human being is at least 18 years of age.

28. The method of claim 27, wherein the human being is at least 40 years of age.

29. The method of claim 27, wherein the human being has a baseline pain intensity score of at least 4 on the 0-10 Numeric Rating Scale.

30. The method of claim 1, wherein the CRPS is CRPS type II.

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